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L1
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               288383-20-0 REGISTRY
RN
ED
               Entered STN: 08 Sep 2000
CN
               Quinazoline, 4-[(4-fluoro-2-methyl-1H-indol-5-yl)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy]-6-methoxy-7-[3-(1-methoxy-7-1)oxy-7-[3-(1-methoxy-7-1)oxy-7-[3-(1-methoxy-7-1)oxy-7-[3-(1-methoxy-7-1)oxy-7-[3-(1-methoxy-7-1)oxy-7-[3-(1-methoxy-7-1)oxy-7-[3-(1-methoxy-7-1)oxy-7-[3-(1-methoxy-7-1)oxy-7-[
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OTHER NAMES:
CN
              4-(4-Fluoro-2-methylindol-5-yloxy)-6-methoxy-7-[3-(pyrrolidin-1-
               yl)propoxy]quinazoline
CN
               4-[(4-Fluoro-2-methyl-1H-indol-5-yl)oxy]-6-methoxy-7-[3-(pyrrolidin-1-indol-5-yl)oxy]
               yl)propoxy]quinazoline
               AZD 2171
CN
              Cediranib
CN
               ZD 2171
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DR
               790713-41-6, 557795-03-6
              C25 H27 F N4 O3
MF
CI
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SR
               CA
LC
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                     TOXCENTER, USAN, USPAT2, USPATFULL
                            (*File contains numerically searchable property data)
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IRINOTECAN-TETRAZOLIUM VIOLET MIXT./CN
IRINOTECAN-VINCRISTINE N-OXIDE MIXT./CN
IRIOBRONZE FINE SILVER/CN
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E12
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=> d
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L2
     97682-44-5 REGISTRY
RN
     Entered STN: 18 Aug 1985
ED
     [1,4'-Bipiperidine]-1'-carboxylic acid,
CN
     (4S)-4,11-diethyl-3,4,12,14-tetrahydro-4-hydroxy-3,14-dioxo-1H-
     pyrano[3',4':6,7]indolizino[1,2-b]quinolin-9-yl ester (CA INDEX NAME)
OTHER CA INDEX NAMES:
     1H-Pyrano[3', 4':6,7]indolizino[1,2-b]quinoline,
     [1,4'-bipiperidine]-1'-carboxylic acid deriv.
CN
     [1,4'-Bipiperidine]-1'-carboxylic acid,
     4,11-diethyl-3,4,12,14-tetrahydro-4-hydroxy-3,14-dioxo-1H-
     pyrano[3',4':6,7]indolizino[1,2-b]quinolin-9-yl ester, (S)-
OTHER NAMES:
CN
   (+)-Irinotecan
CN
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MF
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       HSDB*, IMSDRUGNEWS, IMSPATENTS, IMSPRODUCT, IMSRESEARCH, IPA, MEDLINE,
       MRCK*, PATDPASPC, PROMT, PROUSDDR, PS, RTECS*, SYNTHLINE, TOXCENTER,
       USAN, USPAT2, USPATFULL
         (*File contains numerically searchable property data)
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Absolute stereochemistry. Rotation (+).

100 REFERENCES IN FILE CA (1907 TO DATE)

6 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

3414 REFERENCES IN FILE CA (1907 TO DATE)
72 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
3431 REFERENCES IN FILE CAPLUS (1907 TO DATE)

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ED
     Entered STN: 16 Nov 1984
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CN
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CN
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CN
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     5-Fluoro-2,4(1H,3H)-pyrimidinedione
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                E 5-FLUOROURACIL/CN
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L3
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         21546 L3
L5
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      25144356 PY<2005
L6
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    ANSWER 1 OF 1 CAPLUS COPYRIGHT 2009 ACS on STN
                         2004:965067 CAPLUS <<LOGINID::20091022>>
ACCESSION NUMBER:
                         141:406039
DOCUMENT NUMBER:
TITLE:
                         Combinations for the treatment of diseases involving
                         cell proliferation, migration or apoptosis of myeloma
                         cells, or angiogenesis
                         Hilberg, Frank; Solca, Flavio; Stefanic, Martin
INVENTOR(S):
                         Friedrich; Baum, Anke; Munzert, Gerd; Van Meel,
                         Jacobus C. A.
PATENT ASSIGNEE(S):
                         Boehringer Ingelheim International G.m.b.H., Germany;
                         Boehringer Ingelheim Pharma G.m.b.H. & Co. K.-G.
SOURCE:
                         PCT Int. Appl., 101 pp.
                         CODEN: PIXXD2
DOCUMENT TYPE:
                         Patent
LANGUAGE:
                         English
FAMILY ACC. NUM. COUNT: 2
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21449 REFERENCES IN FILE CA (1907 TO DATE)

500 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA

PATENT INFORMATION:

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KIND DATE APPLICATION NO. DATE
     PATENT NO.
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     WO 2004096224 A2 20041111 WO 2004-EP4363
WO 2004096224 A3 20041216
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              GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK,
              LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO,
              NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ,
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                                20041111 AU 2004-233576
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20060208 EP 2004-729366
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     BR 2004009919 A 20060425 BR 2004-9919
                                                                         20040424
     JP 2006524634 T 20061102 JP 2006-500099
IN 2005DN04018 A 20091002 IN 2005-DN4018
MX 2005011656 A 20051215 MX 2005-11656
NO 2005005605 A 20051128 NO 2005-5605
                                                                         20040424
                                                                         20050907
                                                                         20051028
                                                                         20051128
                                                EP 2003-9587
                                                                    A 20030429
PRIORITY APPLN. INFO.:
                                                EP 2004-508
                                                                    A 20040113
                                                EP 2004-1171
                                                                     A 20040121
                                                WO 2004-EP4363 W 20040424
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AB The present invention relates to a pharmaceutical combination for the treatment of diseases which involves cell proliferation, migration or apoptosis of myeloma cells, or angiogenesis. The invention also relates to a method for the treatment of said diseases, comprising co-administration of effective amts. of specific active compds. and/or co-treatment with radiation therapy, in a ratio which provides an additive and synergistic effect, and to the combined use of these specific compds. and/or radiotherapy for the manufacture of corresponding pharmaceutical combination prepns. The pharmaceutical combination can include selected protein tyrosine kinase receptor antagonists and further chemotherapeutic or naturally occurring semisynthetic or synthetic agents.

IC ICM A61K031-496

=> d his

(FILE 'HOME' ENTERED AT 17:49:55 ON 22 OCT 2009)

FILE 'REGISTRY' ENTERED AT 17:50:43 ON 22 OCT 2009
E CEDIRANIB/CN

L1 1 S E3

FILE 'CAPLUS' ENTERED AT 17:51:19 ON 22 OCT 2009

FILE 'REGISTRY' ENTERED AT 17:51:28 ON 22 OCT 2009 E IRINOTECAN/CN L2 1 S E3 E 5-FLUOROURACIL/CN L3 1 S E3 FILE 'CAPLUS' ENTERED AT 17:52:26 ON 22 OCT 2009 0 S L1 (L) (L2 OR L3) L4L5 26 S L1 AND (L2 OR L3) 1 S L5 AND PY<2005 L6 => s 15 and py=2005 1432022 PY=2005 3 L5 AND PY=2005 => d ibib 1-3 L7 ANSWER 1 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN ACCESSION NUMBER: 2005:1075607 CAPLUS <<LOGINID::20091022>> DOCUMENT NUMBER: 143:339615 TITLE:

AZD2171 in combination with 5-FU and/or CPT-11 for the

treatment of cancer

Wedge, Stephen Robert
Astrazeneca AB, Swed.; Astrazeneca UK Limited
PCT Int. Appl., 47 pp. INVENTOR(S):

PATENT ASSIGNEE(S): SOURCE:

CODEN: PIXXD2

DOCUMENT TYPE: Patent English LANGUAGE:

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

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															FΙ,			
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				•										•	SK,			
															YU,			
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EP	1729808																	
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	2006007555					2008			ZA 2									
	2006010758			A 20061215					MX 2		-							
	20080125447								US 2									
_	2006004755					2006	-		NO 2									
KR	2006	1307	63		A		2006	1219		KR 2	006-	7217	74		2	0061	020	

GB 2004-6446 A 20040323 WO 2005-GB1080 W 20050322 PRIORITY APPLN. INFO.:

REFERENCE COUNT: 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

ANSWER 2 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2005:99470 CAPLUS <<LOGINID::20091022>>

DOCUMENT NUMBER: 142:197889

TITLE: Fluoro substituted omega-carboxyaryl diphenyl urea for

treatment of raf, VEGFR, PDGFR, p38 and flt-3

kinase-mediated diseases

Dumas, Jacques; Boyer, Stephen; Riedl, Bernd; Wilhelm, INVENTOR(S):

Scott

PATENT ASSIGNEE(S): Bayer Pharmaceuticals Corporation, USA

SOURCE: PCT Int. Appl., 68 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent LANGUAGE: English

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

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CN	CN 1856469				A		2006	1101		CN 2	004-	8002		20040722						
JP	JP 2006528196				Τ		2006	1214		JP 2	006-	5212	20040722							
ES	ES 2297490						2008	0501	JP 2006-521221 ES 2004-786091						20040722					
ZA	2006	0006	09		Α		2007	0530		7.A 2	006-	609			2	0060				
KR	2006	0528	66		Α		2006	0519	KR 2006-701558						20060123					
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REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L7 ANSWER 3 OF 3 CAPLUS COPYRIGHT 2009 ACS on STN

ACCESSION NUMBER: 2004:965067 CAPLUS <<LOGINID::20091022>>

DOCUMENT NUMBER: 141:406039

TITLE: Combinations for the treatment of diseases involving

cell proliferation, migration or apoptosis of myeloma

cells, or angiogenesis

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